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PAIN

Fear, boldness and caution: parent effects on how children manage chronic pain --Manuscript Draft--

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Corresponding Author:	Jeremy Gauntlett-Gilbert, Ph.D. Royal United Hospital Bath NHS Trust Bath, UNITED KINGDOM
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Corresponding Author's Secondary Institution:	
First Author:	Jeremy Gauntlett-Gilbert, Ph.D.
First Author Secondary Information:	
Order of Authors:	Jeremy Gauntlett-Gilbert, Ph.D. Abbie Jordan, Ph.D.
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Prof Frank Keefe

Editor, *PAIN*

Dear Prof Keefe,

Re: Invited Commentary on Birnie et al.

Many thanks for the invitation to submit a commentary on this interesting paper. We hope to have drawn out the paper's novel and positive aspects, as well as situating it more broadly in the current literature.

We look forward to hearing your view on this submission.

Yours sincerely,

Jeremy Gauntlett-Gilbert

Abbie Jordan

Fear, boldness and caution: parent effects on how children manage chronic pain

Jeremy Gauntlett-Gilbert^{1,2}*

Abbie Jordan³

1 – Bath Centre for Pain Services, UK

2 – Faculty for Health and Applied Sciences, University of the West of England, UK

3 – Centre for Pain Research & Department of Psychology, University of Bath, UK

- * Corresponding author –

Dr Jeremy Gauntlett-Gilbert, Bath Centre for Pain Services, Royal United Hospitals Bath, Bath
BA1 3NG, UK.

jeremy.gauntlett-gilbert@nhs.net; +44 1225 821181

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4 It is well-established that parents affect how children with chronic pain manage their pain and
5 associated disability, yet exactly how that influence happens is not fully understood. There is
6 substantial evidence for 'intergenerational transmission' of pain and pain related disability(1), with
7 mothers with chronic pain having children at risk of poorer outcomes across a number of
8 domains(2). Studies have also shown that children of parents with a particular type of pain are more
9 likely to develop similar pain, or other pain diagnoses(3).

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15 However, not all children with chronic pain have parents who also experience chronic pain, and it
16 remains important to establish the exact mechanisms of parental influence. The literature on
17 intergenerational transmission has examined a wide range of variables that may pass risk from
18 mother to child(1). However, a number of these mechanisms, such as maternal genetics, prenatal
19 and perinatal influences, are not manipulable in the present moment and consequently, are unable
20 to inform pain treatment. Thus, researchers have worked to establish the role of a number of key
21 parent behavioral variables as potential treatment targets, for example, parent pain
22 catastrophizing(4), pain acceptance (5), and protectiveness(6). However, results have not always
23 been consistent; for example, some studies have shown a role for protectiveness in affecting child
24 pain outcomes(7), whilst other studies have not(8). Birnie and colleagues, in this volume, approach
25 this problem by assuming that the influence of such parent variables may vary in different
26 contexts(9); they may be driven by poor parent health and further moderated by parent pain status
27 and child age. In particular, they argue that research focus should shift from examining parent pain
28 to all forms of parent ill health, which is a novel aspect of this study.

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41 In their study, Birnie et al. use a conceptually clear clinical model, the Interpersonal Fear-Avoidance
42 Model (IFAM)(7) to examine a centrally important clinical outcome, pain impact in the child. The
43 IFAM suggests that important sources of child disability – pain catastrophizing and avoidance in the
44 child, for example – are affected by the parent's attitude and reaction to the child's pain. Where a
45 parent is catastrophic, hypervigilant and overprotective, their child will likely increase their own
46 anxious reactions to pain, to their detriment.

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52 This interaction between parents and children is evidently complex and multifactorial, often
53 examined with structural equation modelling (SEM). The sheer number of potential influences can
54 lead investigators to create elaborate models. In contrast, Birnie et al. have stayed close to a
55 theoretical model, the IFAM, and then used a clear and constrained model. There are relatively few
56 variables and paths, and each piece yields information; multiple group comparisons are used rather

1 than burdening the model with extra variables. Their large dataset, with highly complete data drawn
2 from an adaptive computerised questionnaire system, make this study particularly valuable.

3
4 Birnie et al.'s results add support to the IFAM by showing that parent catastrophizing seems to drive
5 child catastrophizing, which in turn worsens pain impact. Parents' own catastrophising seems to be
6 worse when they themselves report poorer health. The authors summarise their results as showing
7 that parental health has an indirect effect on child pain impact that is mediated by parent
8 catastrophizing. However, despite the authors' interest in parental health, the results also point
9 to the centrality of behavioural and psychological factors. Specifically, parent catastrophizing seems
10 to the key mechanism of negative effects and there is nothing about a parent simply being less
11 healthy, for example, that leads to them acting more protectively towards their child (the latter
12 again being related to parent catastrophizing).

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14 The analysis of child age and parent pain status, as moderators of IFAM mechanisms, yields
15 interesting results. Parent protectiveness was found to be more unhelpful in adolescents, compared
16 to younger children, perhaps due to its limiting influence on adolescent autonomy. However, the
17 actual size of the effects across age was similar, and small. Looking at parent pain status, the
18 negative effects of parent ill health and child catastrophizing were relevant in both groups, but
19 surprisingly stronger where parents did not report pain. This raises the interesting possibility that
20 pain catastrophizing may be more potent, for the parent and the child, when pain is more unfamiliar
21 in the family.

22
23 The results of Birnie and colleagues' study emphasise that even the most powerful and familiar
24 parent variables, such as catastrophizing and protectiveness, may have quite different influences in
25 different family and developmental contexts. Attention to such moderators of parent effects may
26 increase replicability of results in this literature. Consequently, it is critical that future studies of
27 parent influence explore child age as a moderator variable. It may also be that the severity and
28 nature of the child's condition will change parent influence. Parent effects are sometimes studied in
29 diverse and contrasting pediatric pain populations. For example, Cordts et al. recently studied
30 parents of children with an average pain of 4.1/10, where 63% were experiencing headache(8); in
31 contrast, Kemani et al.'s population reported much higher pain at 7.3/10 with <15% with
32 headache(10). It is not clear that such populations are comparable or whether we would predict
33 identical parent effects in both. Inconsistent reporting of variables that characterise the severity or
34 duration of a population's problems can also make it hard to compare research findings. For
35 example, Birnie et al. report pain duration but not pain intensity for their participants, whereas
36 Cordts et al. report the opposite. Future investigators should ideally include details of the child's

pain intensity, duration, and functional impact. Such details, along with moderator analyses of child age and parent characteristics, may help to explain inconsistencies in previous research results.

Conflict of interest statement: The authors have no conflicts of interest to declare.

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